



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<b>(21) International Application Number:</b> PCT/EP98/03824 <b>(22) International Filing Date:</b> 23 June 1998 (23.06.98) <b>(30) Priority Data:</b> 97/08275                      1 July 1997 (01.07.97)                      FR <b>(71) Applicant (for all designated States except CA FR US):</b> SOF-ITECH N.V. [BE/BE]; Rue de Stalle 142, B-1180 Brussels (BE). <b>(71) Applicant (for CA only):</b> SCHLUMBERGER CANADA LIMITED [CA/CA]; 24th floor, Monenco Place, 801 6th Avenue, S.W., Calgary, Alberta T2P 3W2 (CA). <b>(71) Applicant (for FR only):</b> COMPAGNIE DES SERVICES DOWELL SCHLUMBERGER [FR/FR]; 50, avenue Jean-Jaurès, F-92541 Montrouge (FR). <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> BARLET-GOUEDARD, Véronique [FR/FR]; Résidence La Baleine, 2, impasse du Cachalot, F-92290 Châtenay Malabry (FR). MAROY, Pierre [FR/FR]; 2, allée Charles Nicolle, F-78530 Buc (FR).		<b>(74) Agent:</b> MENES, Catherine; Etudes & Productions Schlumberger, Division Dowell, 26, rue de la Cavée, Boîte postale 202, F-92142 Clamart Cedex (FR). <b>(81) Designated States:</b> AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). <b>Published</b> <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
<b>(54) Title:</b> CEMENTING COMPOSITIONS AND APPLICATION THEREOF TO CEMENTING OIL OR ANALOGOUS WELLS <b>(57) Abstract</b> <p>The present invention relates to a cementing composition for an oil or analogous well based on Portland cement and silica. The composition also includes a certain quantity of a mineral oxide suitable for promoting the formation of a phase that is temperature- and pressure-stable, such as a hydrogarnet. The invention is of particular application to cementing wells subjected to high temperatures and which may also be subjected to chemical attack, in particular by brine.</p>		

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## CEMENTING COMPOSITIONS AND APPLICATION THEREOF TO CEMENTING OIL OR ANALOGOUS WELLS

- 5 The present invention relates to techniques for drilling oil, gas, water, geothermal, and analogous wells. More precisely, the invention relates to cementing compositions, particularly those adapted for high temperature wells such as geothermal wells, very deep wells, or more generally wells which suffer thermal attack, in particular by injection of steam to stimulate production and/or subterranean brine attack.
- 10 After drilling an oil or analogous well, a casing or a coiled tube is lowered into the well and is cemented over all or part of its length. Cementing primarily serves to prevent fluid exchange between the various formation layers traversed by the well, to prevent gas from rising via the annulus surrounding the casing, or to limit ingress of water into the production well. Its main purpose, of course, is to ensure that the casing is held in place.
- 15 In the vast majority of cases, Portland cement is used, which is a material that is essentially constituted by calcium silicates which are hydrated to form a hydrated monocalcic silicate and lime. That hydrous silicate, conventionally known as C-S-H, is a gel which is largely responsible for the cohesion and strength of the set cement. When the temperature exceeds about 100°C, that gel metamorphoses into a highly crystalline phase termed alpha dicalcic
- 20 silicate  $\alpha\text{-C}_2\text{SH}$  which is much denser than the C-S-H phase with a contraction in the volume and an increase in the porosity of the cement. This results in more rapid disintegration of the cement as its greater porosity also renders it more sensitive to chemical attack such as brine washing or carbon dioxide circulation.

In order to avoid conversion of C-S-H to  $\alpha\text{-C}_2\text{SH}$ , the lime to silica ratio is normally

25 reduced by adding silica in the form of quartz in a proportion of 35% to 40% relative to the weight of Portland cement. Under those conditions, the gel is transformed into tobermorite  $\text{C}_5\text{S}_6\text{H}_5$  at about 110°C then into xonotlite if the temperature is raised to 150°C. Those two minerals have good mechanical strength, but their durability at high temperature and especially their resistance to chemical attack are relatively low. Degradation of the structure

30 is more rapid when the system has already started to degrade, due in particular to an increase

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in the porosity of the cement. Further, near 250°C, a new mineralogical phase, truscottite ( $C_7S_{12}H_3$ ) appears which further weakens the cement. The various degradation mechanisms mentioned above are not exhaustive, of course, especially since no mention has been made of the role of the other oxides present in Portland cement, in particular  $C_3A$  aluminates, and  
5  $C_4AF$  ferrites, but they are sufficient to show the complexity of the phenomena occurring, even without considering the reaction products between the cementing matrix and brines.

Since the disintegration of cement is partly due to an increase in their porosity, it has been proposed to counter this by using plastics additives such as styrene-butadiene latex. However, such additives are themselves rendered unstable by increasing temperature and  
10 they retard rather than prevent cement degradation.

Such problems of cement disintegration with rising temperature are felt more keenly in the industry with increasing use of very deep wells, of steam injection to recover very viscous oil, or of geothermal wells for the exploitation of geothermal energy.

The aim of the invention is to propose cementing compositions which can withstand high  
15 temperatures, of up to 250°C to 300°C, for example, and which are preferably also capable of resisting chemical attack which is often the consequence in high temperature applications.

The path followed by the inventors was to select a mineralogical phase which is stable at the envisaged temperatures and pressures and to promote its formation by adding a certain quantity of a mineral oxide to a base constituted by Portland cement and silica.

20 In a first embodiment of the invention, the formation of calcic hydrogarnets from the series  $[SiO_4]Ca_3M_2^{3+}$  is promoted, where  $M = Al$  (grossularite) or  $M = Fe$  (andradite) by the addition of alumina or ferrite. The quantity added is typically in the range 20% to 50% (by weight relative to the weight of Portland cement), i.e., of the same order of magnitude as the added silica.

25 It should be noted that patent application SU-A-1 654 540 describes a composition for a geothermal well constituted by Portland cement, quartz sand, and a small quantity of an iron cement, a by-product from steel refining, which is relatively rich in alumina (35% to 37% [percentages by weight]), lime (40%), and silica (7% to 10%). It is indicated that adding iron cement encourages hydrogarnet formation, but it should be pointed out that because of  
30 the very small quantity of aluminium and iron present in the mixture, only a very small

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quantity of that very stable mineralogical phase can form. Further, iron cement also provides large quantities of lime while, as indicated above, the principal aim of adding silica is to reduce the lime to silica ratio. Further, the addition of alumina and iron is effected by silico-aluminates which are considered to be reactive while in the invention, the alumina or  
5 iron oxides added to promote the hydrogarnets are considered to be of low reactivity.

A particularly preferred composition is constituted by class G Portland cement, 30% to 40% of silica (preferably about 35%) and 20% to 40% of alumina (preferably about 30%). The cement obtained is particularly resistant to persistent thermal shocks at high temperatures, of about 300°C, for example. Further, the grossularite hydrogarnets formed at high  
10 temperatures react with brines to produce feldspathoids, i.e., mineral phases which are themselves particularly resistant.

Further details and advantageous characteristics of the invention appear from the following description of tests carried out on different examples of additive compositions, made with reference to the following drawings in which:

- 15 • Figure 1 is a phase diagram of a ternary silica/aluminium/calcium system;
- Figure 2 is a graph showing how compressive strength varies after high temperature treatment, for different cement compositions;
- Figure 3 is a diagram showing the test protocol illustrating the resistance of cements to thermal shock;
- 20 • Figure 4 is a diagram showing the test protocol illustrating the resistance of cements to "chemical" shock.

Figure 1 is a phase diagram showing the principal crystalline phases of hydrates found in a silica/aluminium/calcium system which can be obtained at temperatures which are of the order of 100°C to 400°C, represented by the solid circles. It should be noted that a single  
25 CaO-SiO<sub>2</sub>-H<sub>2</sub>O system can form about twenty hydrates which are recorded in the literature and this diagram is therefore not exhaustive.

With a calcium/silica ratio of the order of 2, class G Portland cement (shown on the figure as the symbol +) is in the hillebrandite C<sub>2</sub>SH stability region which forms at about 150°C and after conversion of the C-S-H gel to  $\alpha$ -C<sub>2</sub>SH, itself a compound which culminates in

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complete disintegration of the cement because of the reduction in the apparent volume of the gel. For this reason, in all applications where the cement is likely to be subjected to temperatures of more than 100°C, silica is added to reduce the calcium/silica ratio.

The symbol ■ corresponds to an addition of 35% of silica (by weight relative to the weight of class G Portland cement) which encourages the formation of tobermorite  $C_5S_6H_5$ , a hydrate which forms when the cement is heated to about 100°C and traces of which are found at temperatures of up to almost 250°C. Tobermorite appears to be encouraged by the presence of aluminium.

If the cement is heated to about 150°C, the hydrate continues to lose its water and the tobermorite is gradually replaced mainly by xonotlite  $C_6S_6H$ , a hydrate which forms more readily when the silica is supplied in the form of quartz. To a far lesser extent, a small amount of gyrolite is also observed to form. These phases exist up to temperatures of about 400°C, but their formation is also accompanied by an increase in the porosity of the cement which weakens its resistance to chemical attack, in particular by brine or carbon dioxide.

The invention aims to modify the reactions in the cement matrix due to an increase in temperature and pressure to produce new mineral phases, in particular hydrogarnets from the hydrogrossularite series, the structure being derived from calcic garnets  $Ca_3Al_2Si_3O_{12}$  ( $C_3AS_3$ ) omitting silicon atoms and replacing oxygen atoms to which they are bonded with hydroxyl groups. When all of the silicon atoms have been replaced,  $C_3AH_6$  is obtained.

As will be shown in the tests described below, this promotion of hydrogarnet phases is obtained if the composition is modified to displace it along the straight line  $\Delta$ , i.e., by adding a certain quantity of alumina. The symbol ○ corresponds to the supplemental addition of 30% of alumina, symbol ☒ to an addition of 50% of alumina. Finally, the symbol □, which is offset from the line  $\Delta$ , corresponds to a formulation on the same base as formulation ○, by adding a further 30% of microsilica.

For these tests, 5 slurries with a slurry density of 1.96 g/cm<sup>3</sup> (16.4 ppg) based on class G Portland cement (Black label) were prepared. The formulations are shown in Table I below. For the liquid constituents (fluid loss control agent and anti-foaming agent), the quantities are shown in gallons per sack of Portland cement (1 gps = 3.78 liters per sack of 94 pounds of cement, i.e., 1 gps = 0.070 liters per kilogram of Portland cement). For all of the other

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constituents, the quantities are in percentages by weight with respect to the weight of Portland cement (BWOC = By Weight of Cement).

All of the proposed formulations were optimized to obtain a slurry with a rheology which was compatible with pumping into an underground well, necessitating the addition of the quantities of dispersing agent shown.

Composition #1 was a prior art petroleum grade cement for high temperature wells, containing 35% of silica. The other compositions contained 20% to 50% of alumina and for composition #4, an additional 30% of microsilica.

<b>TABLE I</b> #	1	2	3	4	5
<b>Al<sub>2</sub>O<sub>3</sub></b>	-	20	30	30	50
<b>Silica</b>	35	35	35	35	35
<b>Micro-silica</b>	-	-	-	30	-
<b>Dispersant</b>	0.3	0.7	0.7	0.7	0.7
<b>Retardant 1</b>	0.32	0.6	0.6	0.3	0.8
<b>Retardant 2</b>	0.16	0.3	0.3	0.15	0.4
<b>Fluid loss control agent</b>	0.6	0.6	0.6	0.6	0.6
<b>Anti-foaming agent 1</b>	-	0.05	0.05	0.05	0.05
<b>Anti-foaming agent 2</b>	0.03	-	-	-	

10

Tables II and III show the results obtained regarding the rheology of the slurry (VP, plastic viscosity and yield point Ty). The measurements were carried out under conditions recommended by the API (American Petroleum Institute), at ambient temperature representative of surface conditions when mixing the cement, and at a temperature of 185°C, i.e., the standard circulation temperature downhole.

15

The values given in Tables II and III indicate that the compositions were well suited to cementing oil or analogous wells.

<b>TABLE II</b> Rheology at mixing temperature (20°C)						
	Plastic viscosity VP		Yield point Ty		Gel strength at 10 min	
#	[cP]	[Pa.s]	[lbf/100ft <sup>2</sup> ]	[Pa]	[lbf/100ft <sup>2</sup> ]	[Pa]
1	602.1	0.602	13.3	6.4	52	24.9
2	145.1	0.145	17.7	8.5	38	18.2
3	100.5	0.100	15.6	7.5	33.5	16.0
4	100.6	0.101	21.7	10.4	37.5	17.9
5	71.25	0.071	20.2	9.7	47	22.5

<b>TABLE III</b> Rheology after conditioning for 20 minutes at 85°C						
	Plastic viscosity VP		Yield point Ty		Gel strength at 10 min	
#	[cP]	[Pa.s]	[lbf/100ft <sup>2</sup> ]	[Pa]	[lbf/100ft <sup>2</sup> ]	[Pa]
1	196	0.196	4.12	1.97	12	5.75
2	61.4	0.061	10.1	4.84	22.5	10.77
3	45.15	0.045	8.7	4.17	19	9.10
4	55.26	0.055	10.2	4.88	37	17.72
5	36.66	0.036	11.7	5.60	27	12.93

Table IV below shows characteristic data for setting cement, in particular its thickening time and compressive strength.

- 5 The thickening time is a measurement made, under dynamic conditions, of the time required for the consistency of a slurry to reach 100 BC measured in standardized units. It should be noted that in practice, a slurry with a consistency of more than 70 BC can no longer be pumped without taking special precautions and in order to avoid major accidents, it is normal to pump only slurries with a consistency of less than 30 BC.
- 10 Estimates of the rate of development of compressive strength were obtained by measuring the rate of propagation of ultrasound through a sample of setting cement under static conditions at 300°F (148.9°C) (measured using an "Ultrasonic Cement Analyser").



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<b>TABLE IV</b> #	1	2	3	4	5
Free water (ml)	1	0	0	0	1
Fluid loss at 300°F (m./30 min)	30	not measured	46	not measured	not measured
Thickening time	3h00	3h39	3h10	3h23	3h29
Compressive strength					
... 3 days at 300°F (148.9°C) psi	7950	5790	4896	6118	not measured
N/cm <sup>2</sup>	5482	3992	3376	4218	
... 1 month at 550°F (287.8°C) psi	5958	6715	7591	3479	2062
N/cm <sup>2</sup>	4108	4630	5234	2399	1422
Deformation (10 <sup>-3</sup> mm/kN)	3.01	not measured	3.40	3.47	not measured
Estimate of compressive strength development					
... 50 psi [34 N/cm <sup>2</sup> ]	3h50	4h53	5h15	3h46	7h09
... 500 psi [848 N/cm <sup>2</sup> ]	4h38	6h53	9h09	4h23	10h51
Compressive strength at 24 h, psi	5028	3603	3440	3309	2101
N/cm <sup>2</sup>	3467	2484	2372	2282	1449

The tested formulations were perfectly stable and no tendency to sedimentation was observed, as shown by the free water volumes.

- 5 Figure 2 shows the compressive strength after 3 days at 148.9°C - 300°F (horizontal hatching) and 1 month at 287.8°C - 550°C (vertical hatching).

At a temperature of about 150°C, after 3 days, the preferred compositions of the invention (#2 and #3) had a compressive strength of the order of 3500 N/cm<sup>2</sup> which is very generally considered to be adequate even if this value is substantially lower than that obtained with  
 10 prior art composition #1. In contrast, if the cement undergoes prolonged thermal attack, it has been shown that rather than decreasing, its compressive strength increases.

Compositions #5 comprising 50% of alumina had a marked retardation in the development of compressive strength and thus are not preferred.

The mechanical properties of composition #4, which in addition to 30% of alumina  
 15 comprised 35% of microsilica, declined even more rapidly than those of reference composition #1.

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Samples were also examined using X rays to study their mineralogical structure and the effect of a thermal shock in water or brine was qualitatively tested.

### **Thermal shock**

- 5 The thermal test was carried out on samples which were allowed to set for 3 days at 300°F (148.9°C). The operating protocol is shown in Figure 3; cement setting and testing were carried out in an autoclave, at a pressure of 3000 psi (208 bars). The crystalline phases were tentatively identified as corresponding to the most probable phases taking into account the micro-analysis and the geometric structure of the observed structures.
- 10 After setting for three days, conventional slurry #1 was seen under the electron microscope as a mass of circular vacuoles resembling geodes filled with hexagonal plates. Crystals of  $C_2SH$ , a little as-yet unreacted silica and a little "tobermorite" were identified. The apparent composition remained the same for reference samples left at 300°F in water for 9 days but an increase in the number of amorphous compounds and a reduction in porosity were
- 15 observed. For samples placed at the same temperature but in a brine (water saturated with sodium sulfate), the crystalline compounds identified remained the same but the matrix became essentially constituted by amorphous structures with small plates dispersed in the matrix. There were no visible alterations on a macroscopic scale.

- In contrast, samples which underwent a thermal shock (9 days at 550°F in water or a brine)
- 20 crumbled merely on contact. In water, the formation of large numbers of needles, identified as xonotlite, was observed. In brine, the presence of pseudocubic crystals was also observed. In both cases, silica and the  $C_2SH$  phase had apparently reacted completely.

- After setting for three days at 300°F, samples #2 and #3 containing respectively 20% and 30% of alumina appeared to be very similar to prior art samples, apart from the presence of
- 25 alumina crystals, and a less porous structure for samples containing the most alumina. At this temperature, a period of 9 days in water or brine altered the structure only a little.

After a thermal shock at 550°F in water, the formation of xonotlite and hydrogarnets was observed, the latter being clearly predominant for samples containing the most alumina. After 9 days in a brine at 550°F, there was observed a clear reduction (sample #2) or even

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almost complete disappearance of the xonotlite (sample #3) accompanied by the formation of a reaction product with the brine, in this case a haüyne type feldspathoid  $[\text{SiAlO}_4]_6(\text{SO}_4)_{2-1}(\text{Na,Ca})_{5-4}$ . In contrast to the prior art samples, the two samples including added alumina were intact.

- 5 The formulations comprising 20% and 30% of alumina were perfectly resistant to thermal shocks in the presence or absence of brine.

### **"Chemical" shock**

10 The "chemical" shock test was carried out on samples left for one month at 500°F (287.8°C) then placed for 9 days, at the same temperature, in a bath of water or of brine. The operating protocol is shown in Figure 4 and as before, cement setting and tests were carried out in an autoclave pressurized to 3000 psi (208 bars).

15 For the conventional cement at the start of the test, the sample principally comprised xonotlite, and also amorphous compounds. Criss-crossed needles were observed which disappeared when the sample was placed in a brine bath to be replaced by small hexagonal prisms.

With cements containing 20% of alumina, the matrix was completely crystallized, with hydrogarnet formation, with no significant evolution after the "chemical" shock.

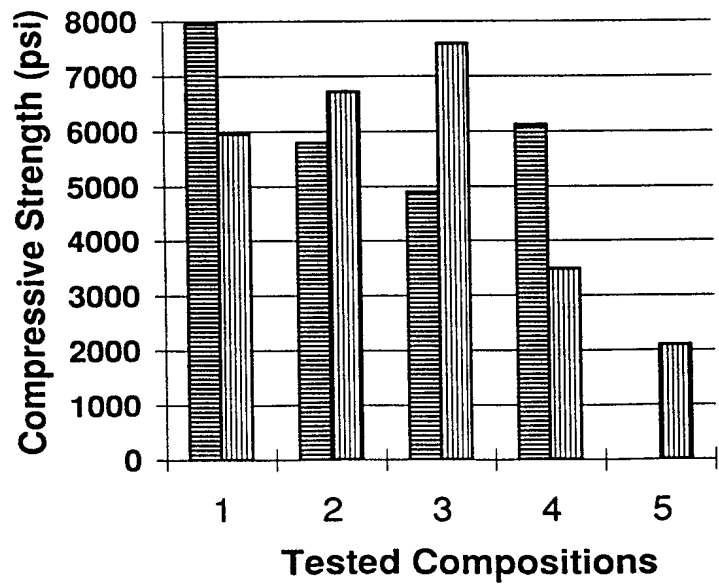
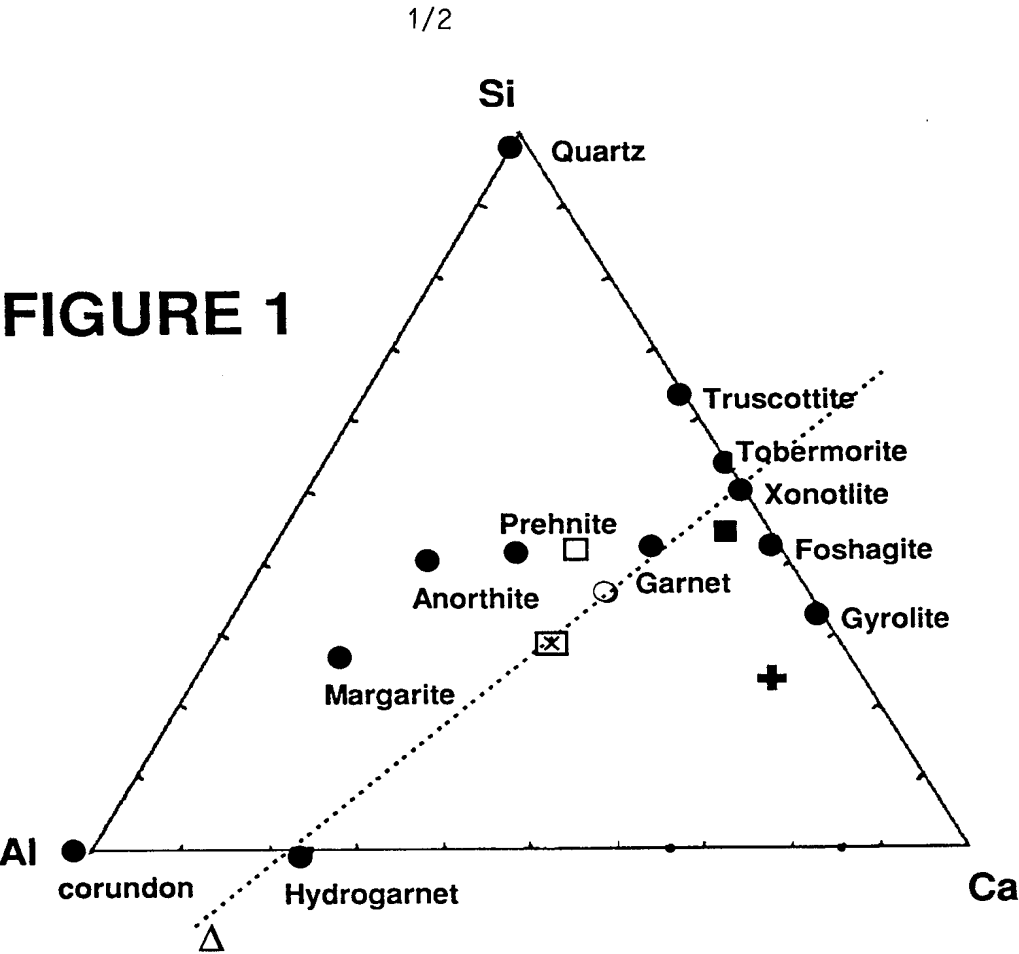
20 With a cement which was richer in alumina, it was observed that the alumina had apparently not yet reacted after one month at 550°F, with the alumina oxides disappearing in samples placed in a water bath. In both cases, the structure was a honeycomb structure with the mesh formed by hydrogarnets filled with the remaining fibers of xonotlite, which structure could explain the remarkable compressive strength of these cements.

25 The formation of a feldspathoid, haüyne, was observed for samples undergoing a chemical shock.

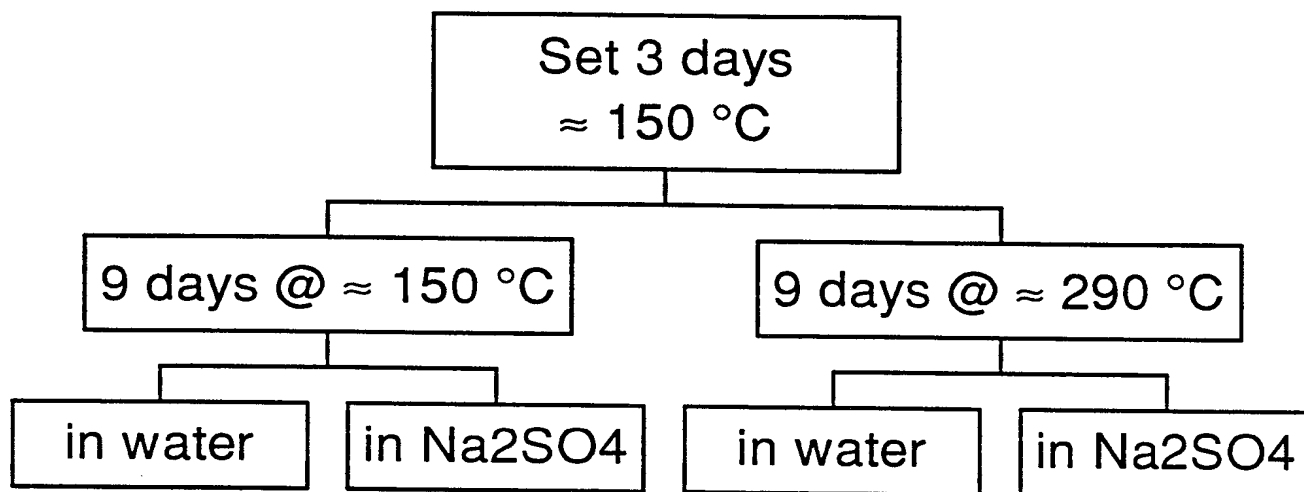
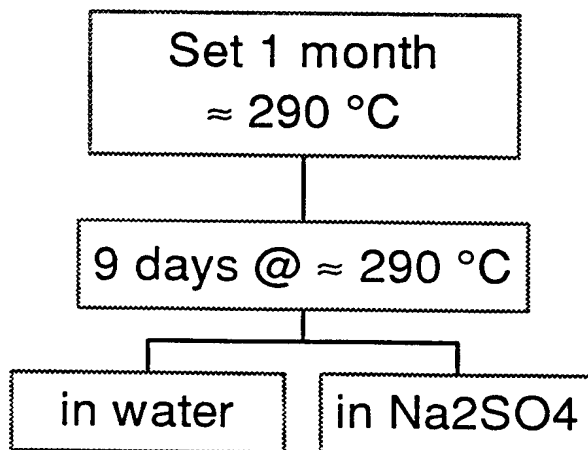
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## CLAIMS

1. A cementing composition for an oil or analogous well, based on Portland cement and  
5 silica, characterized in that it further includes a certain quantity of a mineral oxide  
suitable for promoting the formation of a phase that is temperature- and pressure-  
stable.
2. A composition according to claim 1, characterized in that said oxide is alumina or  
ferrite and encourages the formation of calcic hydrogarnets from the series  
10  $[\text{SiO}_4]\text{Ca}_3\text{M}^{3+}$ , with  $\text{M}=\text{Al}$  (grossularite) or  $\text{M}=\text{Fe}$  (andradite).
3. A composition according to claim 2, characterized in that it includes alumina in a  
proportion of 20% to 50% (by weight relative to the weight of Portland cement).
4. A composition according to claim 2, characterized in that it is constituted by  
Portland cement, silica (30% to 40% by weight relative to the weight of Portland  
15 cement), and alumina (30% to 40% by weight relative to the weight of Portland  
cement).
5. Application of compositions according to any one of claims 1 to 3 to cementing  
wells which are continuously or repeatedly subjected to temperatures of 250°C to  
300°C.



**FIGURE 2**

**FIGURE 3****FIGURE 4**

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 98/03824

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC 6 C04B22/06 E21B33/13

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

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Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>DATABASE WPI  Week 9220  Derwent Publications Ltd., London, GB;  AN 92-164729  XP002060045  &amp; SU 1 654 540 A (BOREHOLE CONSOLIDATION  MUDS)  cited in the application  see abstract</p> <p style="text-align: center;">--- -/--</p>	1, 2, 5



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

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Date of the actual completion of the international search

26 October 1998

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Daeleman, P

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 98/03824

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	CHEMICAL ABSTRACTS, vol. 107, no. 12, 21 September 1987 Columbus, Ohio, US; abstract no. 101684j, T. NAGAFUCHI ET AL.: page 322; XP002060043 see abstract & JP 61 286252 A (ID.) ---	1,2
A	CHEMICAL ABSTRACTS, vol. 92, no. 26, 30 June 1980 Columbus, Ohio, US; abstract no. 219972b, O. NAKAMURA: page 281; XP002060044 see abstract & JP 55 003362 A (ID.) ---	1,2
A	GB 2 063 240 A (LAFARGE) 3 June 1981 see page 1, line 36-43 ---	1-3
A	DE 12 63 572 B (H. HARTMANN) 14 March 1968 see claim 1 ---	1-4
A	G.N. CHANGIL'DIN ET AL.: "Ton-Zement-Mörtel zur Zementierung von Gas- und Erdölbohrungen" CHEMISCHES ZENTRALBLATT, no. 11, 1968, page 247 XP002060042 Abrégé nr. 11-2300 -----	1,2,5



# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 98/03824

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
GB 2063240	A	03-06-1981	LU 81915 A	04-06-1981
			AU 536984 B	31-05-1984
			AU 6415180 A	28-05-1981
			BE 886239 A	18-05-1981
			BR 8007555 A	02-06-1981
			CA 1138903 A	04-01-1983
			CS 8007829 A	13-06-1985
			DE 3041652 A	27-05-1981
			DK 487380 A	20-05-1981
			FR 2470103 A	29-05-1981
			IE 50476 B	30-04-1986
			IN 152460 A	21-01-1984
			JP 56145140 A	11-11-1981
			NL 8006298 A	16-06-1981
			OA 6655 A	30-09-1981
			SE 448450 B	23-02-1987
			SE 8008088 A	20-05-1981
			ZA 8006783 A	28-10-1981
<hr/>				
DE 1263572	B		NONE	
<hr/>				